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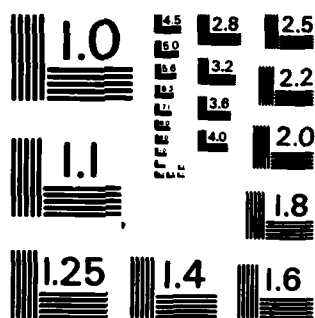
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POINT-BY-POINT MATRIX EFFECT CALIBRATION
FOR THE QUANTITATIVE ANALYSIS OF LAYERED SEMICONDUCTORS
BY SECONDARY ION MASS SPECTROMETRY

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BRIEF

Using the linear dependence of secondary ion yields and sputtering yields on matrix composition, a point-by-point matrix effect calibration is applied to $\text{Al}_x\text{Ga}_{1-x}\text{As}$ multilayer-multimatrix samples.

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ABSTRACT

Point-by-point matrix effect calibration is applied to a variety of $\text{Al}_x\text{Ga}_{1-x}\text{As}$ multilayer-multimatrix samples grown by molecular beam epitaxy. The procedure uses the linear dependence of secondary ion yields and sputtering yields on matrix composition to quantify depth profiles through matrix gradients and interfaces. The proposed method provides accurate results in the analysis of samples too complex for conventional quantitative analysis by secondary ion mass spectrometry.

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Secondary ion mass spectrometry (SIMS) is often used to monitor elemental depth distributions in solids and solid interfaces. The technique is highly sensitive for most elements and has good depth resolution. However, the complexity of the sputtering event has made quantitative analysis difficult. Ion implant standards (1-3) have been successfully used to calibrate the depth profiles of trace elements in homogeneous matrices. However, due to the variation of secondary ion yields and sputtering yields with matrix composition, matrix effects, the quantification of SIMS profiles in multimatrix samples remains a problem.

For $\text{Al}_x\text{Ga}_{1-x}\text{As}$ and related matrices, it has recently been shown that practical ion yields τ (ions detected/atoms sputtered) and sputtering yields S (secondary atoms/primary ion) vary linearly with sample composition (4). In addition, highly precise calibration lines were obtained using relative ion yields $R\tau$ and relative sputtering yields RS . These relative values were obtained by normalizing ion yields and sputtering yields from a sample matrix (τ_x and S_x) to those from a standard matrix (τ_0 and S_0) when both measurements were performed under near identical analysis conditions.

$$R\tau = \tau_x / \tau_0 \quad (1)$$

$$RS = S_x / S_0 \quad (2)$$

In this investigation, the application of these calibration lines to the analysis of $\text{Al}_x\text{Ga}_{1-x}\text{As}$ multilayer-multimatrix samples is examined. These *superlattices*, as shown in Figure 1, are best characterized as a series of homogeneous matrices. A profile correction program (SLIC), which treats each point of a depth profile as a homogeneous matrix, will be presented. SLIC determines the matrix composition at each point of a depth profile and subsequently performs a point-by-point correction of the trace element distributions. The capabilities and limitations of this method will be discussed.

EXPERIMENTAL SECTION

Sample Preparation. The $\text{Al}_x\text{Ga}_{1-x}\text{As}$ matrices were grown by molecular beam epitaxy (MBE) on semi-insulating GaAs substrates. The matrix compositions were determined from the MBE growth parameters and verified to an accuracy of better than 10% (5) using Rutherford backscattering spectroscopy (RBS). In some instances dopants were introduced during the growth process while in others, including the standards, ion implantation was used. Prior to implantation samples were cleaned with acetone and methanol.

Instrumentation. The $\text{Al}_x\text{Ga}_{1-x}\text{As}$ layers were grown in a VARIAN MBE-360 machine (6). RBS measurements were carried out on a GENERAL IONEX Tandetron Model 4110A. Analyses were performed using a 2.7 Mev

He^+ ion beam with solid state detection at a 170° angle from the incident beam path. Ion implantation was performed using a hot filament ion source and a magnet for mass separation.

SIMS analysis was carried out on a CAMECA IMS-3F ion microanalyzer (7) interfaced to a HEWLETT PACKARD 9845T microcomputer for control and data acquisition. A $1.0 \mu\text{A O}_2^+$ primary beam at an energy of 5.5 KeV was rastered over a $300 \times 300 \mu\text{m}$ area. Positive secondary ions were monitored from an image field $60 \mu\text{m}$ in diameter. Analyses were performed with a residual pressure of 2×10^{-8} torr and an energy window of 130 eV. A multiple sample holder was used to simultaneously mount several samples. Depth measurements on the sputtered craters were performed on a TALYSTEP stylus device.

Software. Programs for instrumental control, data analysis, and matrix correction were written in BASIC for the HP 9845T. The program SLIC (superlatttice and interface calibration) was used to correct depth profiles for matrix effects.

SLIC is a comprehensive matrix correction program. For $\text{Al}_x\text{Ga}_{1-x}\text{As}$ matrices, the matrix composition and depth at each point of a depth profile is determined by an iterative process involving calibration lines for both relative sputtering yield and the relative ion yield of $^{75}\text{As}^+$. Dopant profiles are then corrected for matrix changes using the appropriate dopant calibration lines.

Procedure. Each sample was mounted with three standards, created by ion implantation, using a multiple sample holder. The standards, including in each case GaAs and two different $\text{Al}_x\text{Ga}_{1-x}\text{As}$ matrices, and the sample were inserted simultaneously to insure nearly identical analysis conditions. After allowing the pressure in the sample chamber to reach a steady-state condition, the primary beam was focused to a spot of about 100 μm in diameter, and the proper mass settings were determined. Standards and sample were analyzed consecutively without changing any instrumental parameters. Using the depth profiles of the standards, matrix calibration lines were constructed and used to correct the sample depth profiles.

RESULTS AND DISCUSSION

R_r and R_s calibration lines can be used to quantify concentration and depth respectively (4). Using Eq. (3), the concentration of analyte at each point of a depth profile C_p (atoms/ cm^3) can be determined.

$$C_p = I_p / R_{r_x} \cdot \tau_0 \cdot D_p \cdot A \quad (3)$$

R_{r_x} is the relative sputtering yield determined from a calibration

line for the appropriate value of x , and τ_0 is the practical ion yield of the analyte in the standard matrix (GaAs). I_p (counts) and D_p (cm) are the signal and depth increment associated with a given point of a depth profile.

Similarly, the erosion rate at a given point of a depth profile \dot{z}_p (cm/sec) can be determined using Eq. (4).

$$\dot{z}_p = RS_x \cdot \dot{z}_0 \cdot N_0 / N_x \quad (4)$$

RS_x is the relative sputtering yield determined from a calibration line for the appropriate value of x , and \dot{z}_0 (cm/sec) is the erosion rate of the standard matrix (GaAs). N_0 and N_x are the atomic densities of the standard and sample matrix respectively.

With Eqs. (3) and (4) a multilayer-multimatrix sample can now be analyzed. The most critical ingredient of such an analysis is the determination of the matrix structure. The program SLIC exploits the fact that the concentration of As in $Al_xGa_{1-x}As$ matrices is a constant, and that the relative ion yield of $^{75}As^+$ in $Al_xGa_{1-x}As$ matrices can be readily calibrated. If the value of $R\tau_x$ in Eq. (3) is expressed in terms of the equation for the $^{75}As^+$ calibration line ($R\tau_x^{As} = x \cdot M_{R\tau}^{As} + 1$; where $M_{R\tau}^{As}$ is the slope of the line), the equation can be manipulated to the following form:

$$x = ((I_p^{As}/C_p^{As} \cdot A \cdot \tau_0^{As} \cdot D_p) - 1) / M_{R\tau}^{As} \quad (5)$$

Except for D_p , the values for all the variables on the right side of Eq. (5) can be readily determined. Since D_p is directly related to \dot{z}_p ($D_p = \dot{z}_p \cdot T_p$; where T_p is the number of seconds per point), it can be expressed in terms of Eq. (4). If ES_x is replaced with the corresponding equation for the RS calibration line, Eq. (6) is obtained.

$$\begin{aligned} D_p &= (M_{RS} \cdot x + 1) \cdot \dot{z}_0 \cdot N_0 / N_x \cdot T_p \\ &= (M_{Rz} \cdot x + 1) \cdot \dot{z}_0 \cdot T_p \end{aligned} \quad (6)$$

M_{RS} is the slope of the RS calibration line, and M_{Rz} is the slope of the corresponding relative erosion rate calibration line ($M_{Rz} = M_{RS} \cdot N_x / N_0$). Once again the values of D_p and x are the only unknown quantities.

For each point of a depth profile, the program SLIC determines the matrix structure by performing the following process. It initially assumes that $x = 0$. The corresponding value of D_p is determined from Eq. (6), and employed in Eq. (5) to obtain a better approximation of x . This process is reiterated until the value of x

converges. In this manner, the matrix structure can be determined despite matrix gradients, interfaces, and plateaus.

Capabilities and Limitations. As shown in Figure 2a, matrix effects can drastically distort SIMS analyses through multilayer-multimatrix samples. This depth profile of a boron implant through a GaAs and a $\text{Al}_x\text{Ga}_{1-x}\text{As}$ layer shows a large irregularity at 31 time units. This irregularity is due to the changing matrix effects at the interface. In Figure 2b, SLIC was used to determine the Al distribution and correct the $^{11}\text{B}^+$ implant distribution. Upon correction, the implant distribution was transformed into the near gaussian shape which was expected.

The crucial factor influencing the quality of the matrix corrections performed by SLIC is the accuracy with which the matrix structure can be determined. This accuracy can be checked with RBS. RBS analysis is accurate to about 10%, and has an detection limit of 1% for $\text{Al}_x\text{Ga}_{1-x}\text{As}$ matrices. In Figure 3, a complex $\text{Al}_x\text{Ga}_{1-x}\text{As}$ sample was analyzed by both SIMS and RBS. The Al concentration at specific regions of the superlattice was determined using SLIC and standard RBS techniques (5). As shown in Table I, the values determined by the two techniques agree quite well. In fact, the two sets of data are not statistically different at a 95% confidence level.

In addition to matrix composition, the point-by-point correlation between matrix structure and dopant distribution is critical. Small

differences between the actual and the measured matrix structure can significantly influence the corrections performed on dopant distributions. As apparent in Figure 4, the $^{75}\text{As}^+$ signal tracks the Al distribution quite well. SLIC uses this correlation to precisely determine matrix structures. However, errors can still result from the correlation between matrix structure and dopant distribution. For example, there is a dead time between the measurement of the matrix signal and the dopant signal for each point of a depth profile. When this dead time is large compared to the abruptness of the matrix changes, the matrix and dopant signals will be obtained from different matrix regions. Consequently, the dopant signal will be calibrated for the wrong matrix composition. Fortunately, this type of error can usually be avoided by minimizing the dead time and reducing the sputtering rate.

Another obstacle to quantitative multilayer-multimatrix analysis concerns molecular interferences. Molecular ions are frequently the source of the high background signals commonly observed during elemental analysis. Molecular signals, like elemental ion yields, can change with matrix composition. When analyzing a multimatrix sample, failure to correct for the changing abundance of interfering molecular ions can lead to erroneous elemental distributions. For example, in Figure 5a, the $^{28}\text{Si}^+$ profile (dotted line) tracks the Al^{2+} profile (solid line) through the alternating GaAs and $\text{Al}_x\text{Ga}_{1-x}\text{As}$ layers even though this region has not been doped with Si. Fortunately, these changing background levels can often be linearly correlated to the

matrix composition. For example, provided the background signals are normalized to erosion rate, a precise background signal calibration line can be obtained for Si in $\text{Al}_x\text{Ga}_{1-x}\text{As}$. The background signals can then be subtracted point-by-point prior to quantification. In Figure 5b, both a background corrected (dotted line) and an uncorrected (dashed line) Si concentration depth profile are presented. In the corrected version, nearly all the distortions produced by the changing background levels are removed. The doped region at the surface remains followed by a region with the Si concentration below the detection limit (5×10^{17} atom/cm³) of Si in GaAs.

The full utility of SLIC can be appreciated when very complex samples are analyzed. In Figure 6, SLIC has been used to quantitatively analyze a complex $\text{Al}_x\text{Ga}_{1-x}\text{As}$ superlattice grown by MBE with Be and Si dopants. In the uncorrected profile, Figure 6a, both the Be and Si distributions follow the $^{75}\text{As}^+$ signal due to the changing matrix effects. Upon calibration, both the Be and Si distributions have changed substantially. As expected from the growth conditions, the Be concentration generally increases as the Al concentration decreases and vice versa. In addition, excluding the surface build up, the Si distribution has generally leveled out at 4×10^{18} atom/cm³. Without SLIC, this type of analysis could not have been made.

In summary, SLIC is a very precise program for applying calibration lines to the problem of matrix effects in complex samples.

Although most of the work to date has been applied to $\text{Al}_x\text{Ga}_{1-x}\text{As}$ matrices, this procedure can be used in related matrices, such as $\text{In}_x\text{Ga}_{1-x}\text{As}$ and $\text{GaAs}_{1-x}\text{Sb}_x$. In addition, work in this laboratory indicates that similar methods can be applied to Group III and V compound matrices in which several elements are changing simultaneously.

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CREDIT

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Table I. Point-by-point Comparison of Al Concentrations
Determined by SIMS and RBS Analyses

Al Concentration ($\times 10^{21}$ atom/cm ³)			
Point	RBS	SIMS	Deviation
1	12.0	13.0	-1.0
2	4.2	3.9	+0.3
3	5.1	4.8	+0.3
4	5.1	4.7	+0.4
5	5.2	4.7	+0.5
6	11.0	11.0	0.0
7	1.0	1.3	-0.3
8	4.9	5.7	-0.8

Avg Deviation = -0.075

FIGURE CAPTIONS

- Figure 1. A hypothetical $\text{Al}_x\text{Ga}_{1-x}\text{As}$ superlattice. The thickness T of the layers can vary from several angstroms to several microns while x can be varied from 0 to 1.
- Figure 2. SIMS depth profile of a 250 Kev $^{11}\text{B}^+$ implant into a $\text{GaAs}/\text{Al}_{0.3}\text{Ga}_{0.7}\text{As}/\text{GaAs}$ sample. (a) uncorrected profiles of $^{11}\text{B}^+$ (...) and $^{75}\text{As}^+$ (—); (b) concentration profiles of B (...) (2.0×10^{18} atom/cm³ full scale) and Al (—) (1.0×10^{22} atom/cm³ full scale).
- Figure 3. SIMS and RBS depth profiles of an $\text{Al}_x\text{Ga}_{1-x}\text{As}$ superlattice. (a) an uncorrected $^{75}\text{As}^+$ SIMS profile; (b) an RBS profile of the total counts from Ga and As.
- Figure 4. An uncorrected SIMS depth profile of $^{75}\text{As}^+$ (—) and $^{27}\text{Al}^{2+}$ (- - -) in an $\text{Al}_x\text{Ga}_{1-x}\text{As}$ superlattice.
- Figure 5. SIMS depth profile of an $(\text{GaAs}/\text{Al}_{0.35}\text{Ga}_{0.65}\text{As})_n$ superlattice doped in the first 0.4 μm with Si. (a) uncorrected profiles of Al^{2+} (—) and $^{28}\text{Si}^+$ (...); (b) concentration (atom/cm³) profiles of Al^{2+} (—), and Si with (...) and without (- - -) background correction.

Figure 6. SIMS depth profile of an $\text{Al}_x\text{Ga}_{1-x}\text{As}$ superlattice doped with Be and Si. (a) uncorrected profiles of $^9\text{Be}^+$ (---), $^{28}\text{Si}^+$ (...), and $^{75}\text{As}^+$ (—); (b) concentration profiles of Be (---) (5.0×10^{19} atom/cm³ full scale), Si (...) (9.0×10^{19} atom/cm³ full scale), and Al (—) (1.5×10^{22} atom/cm³ full scale).

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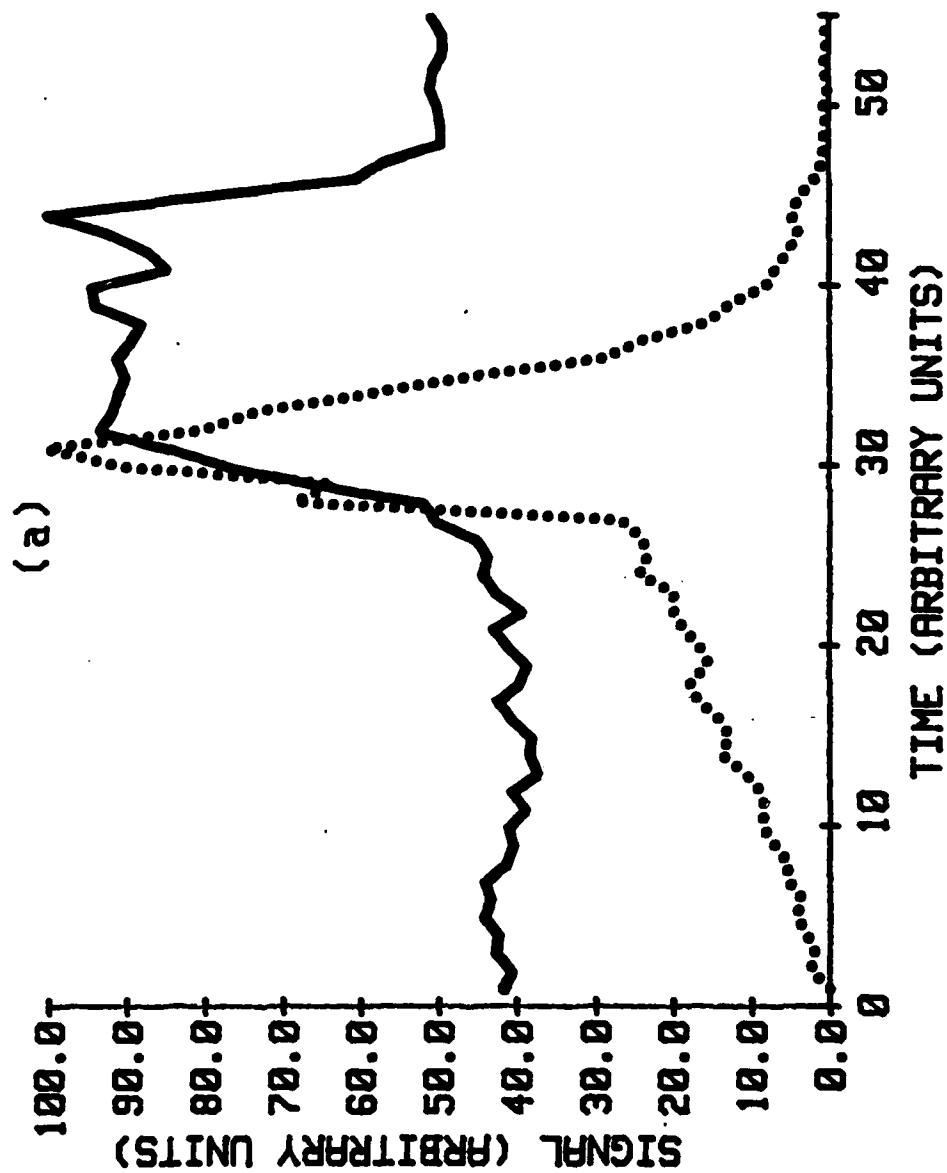
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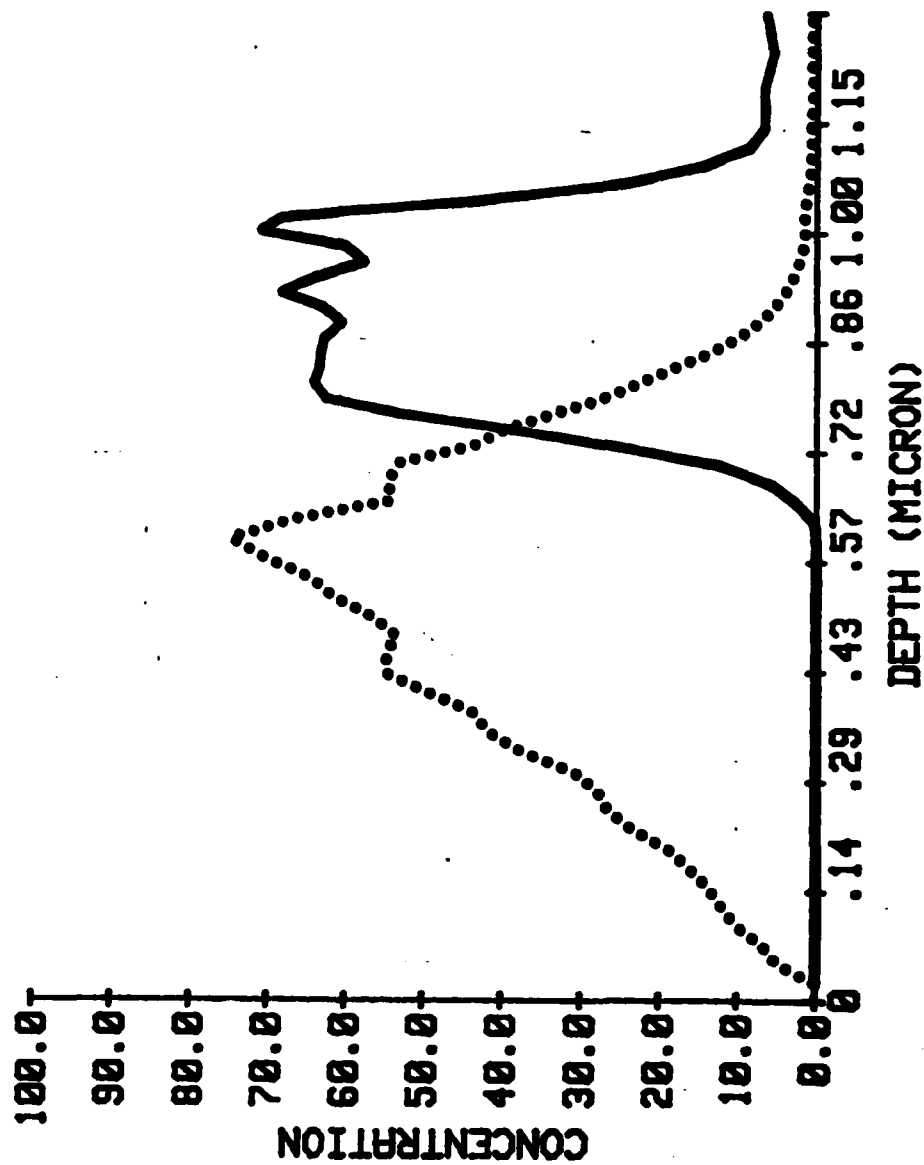
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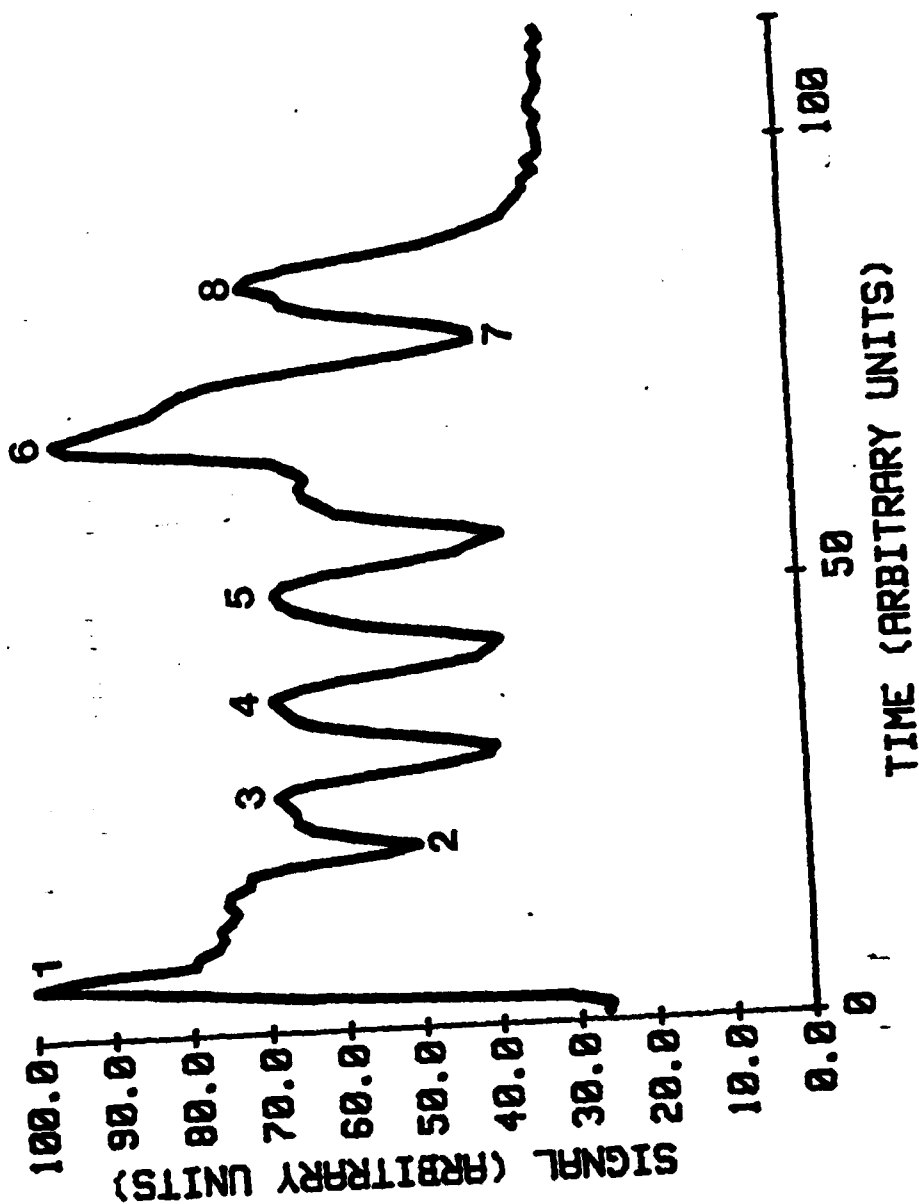
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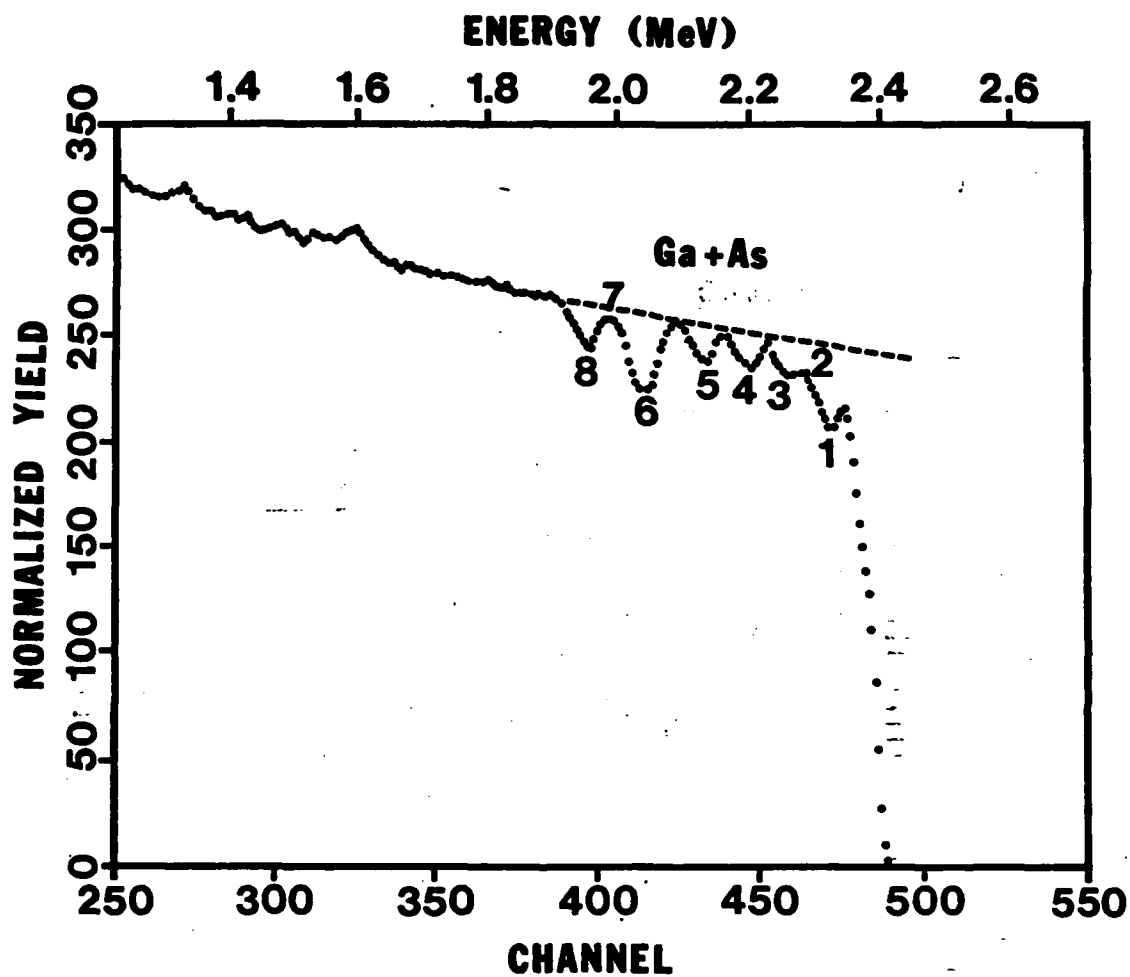
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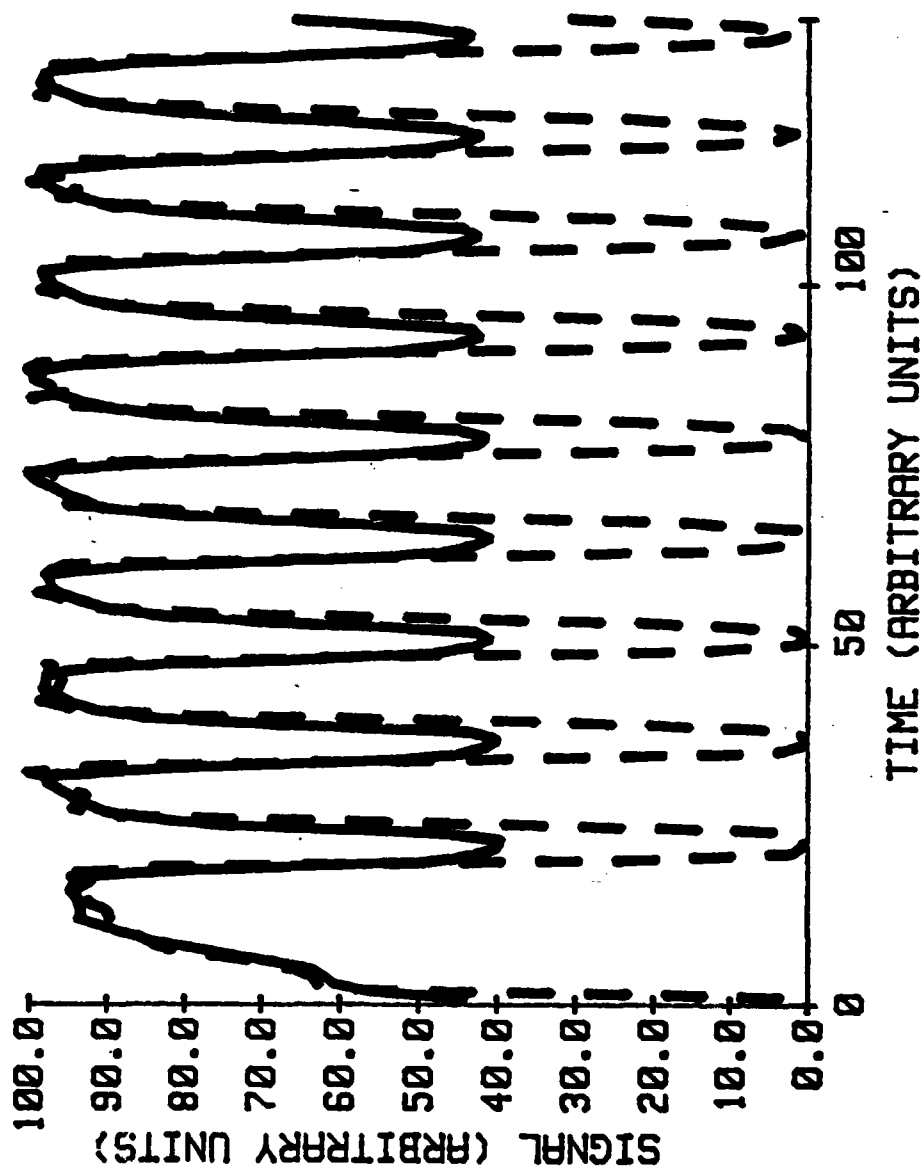


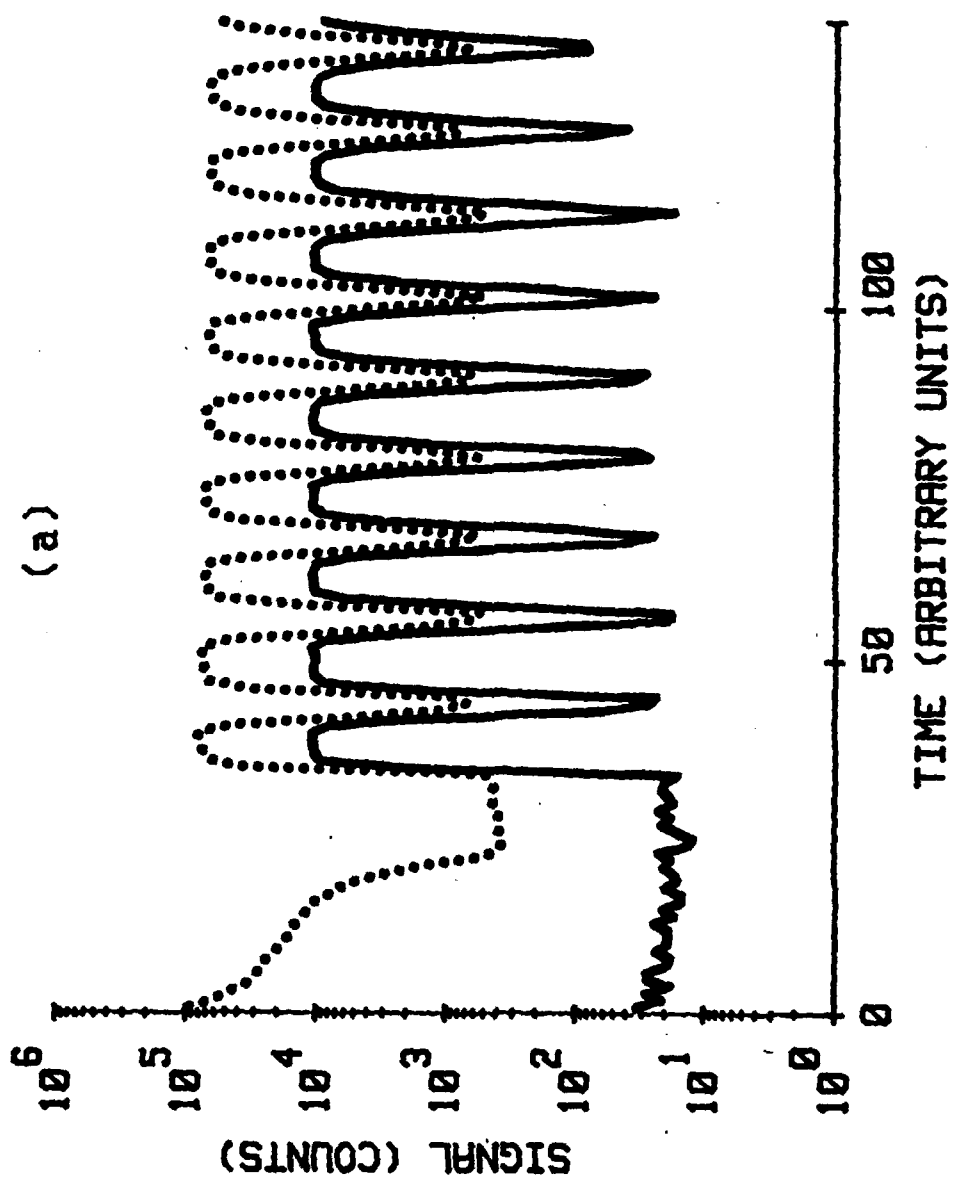
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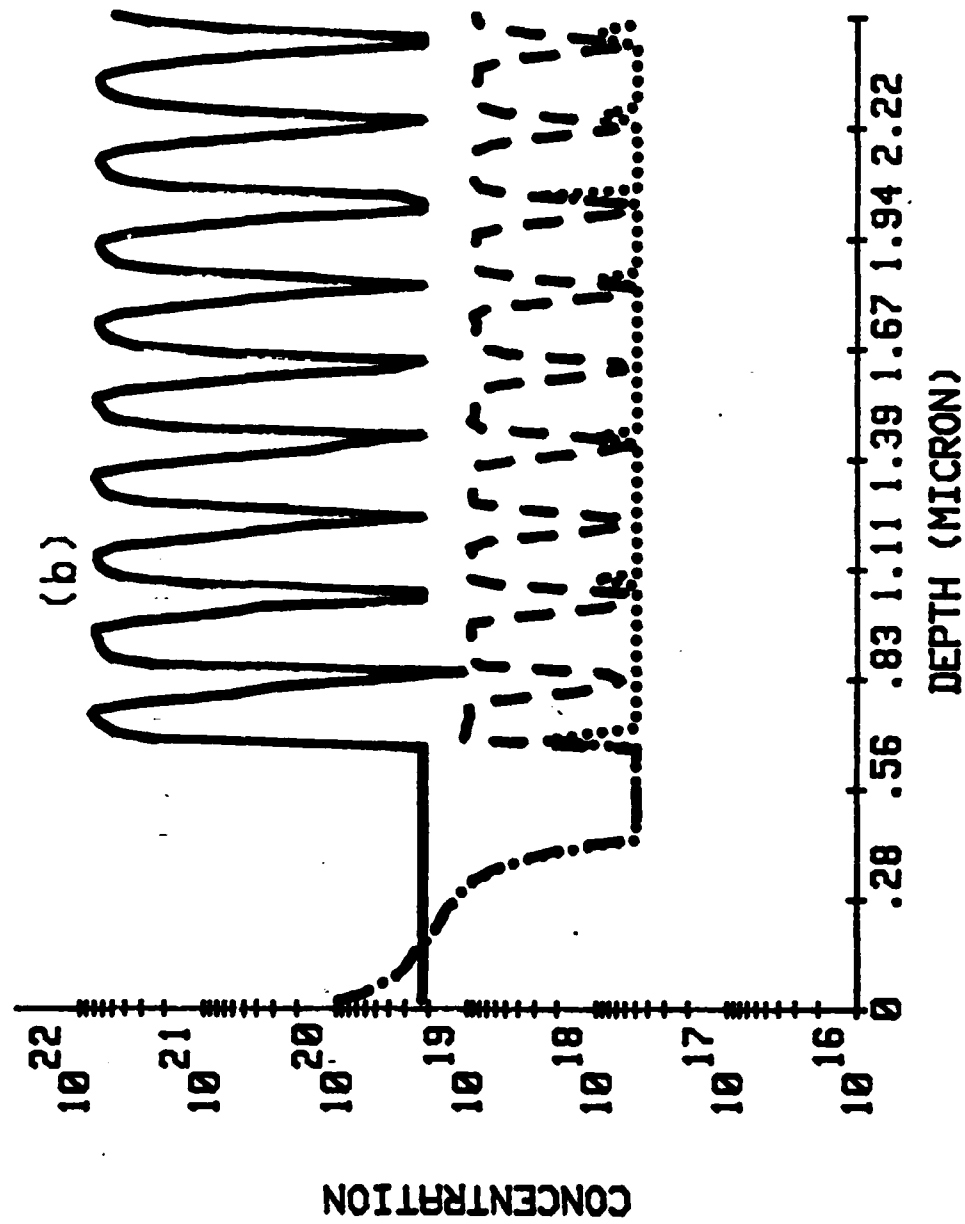


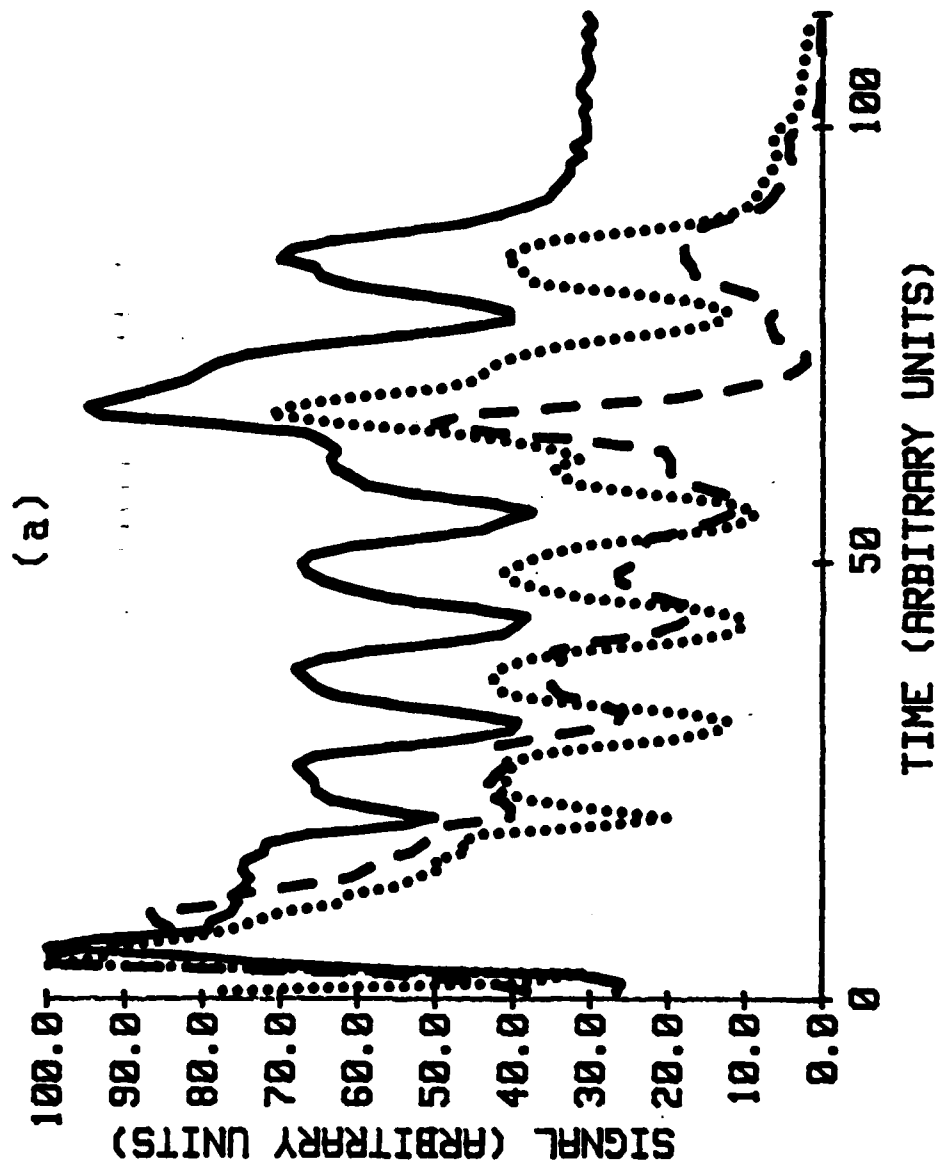
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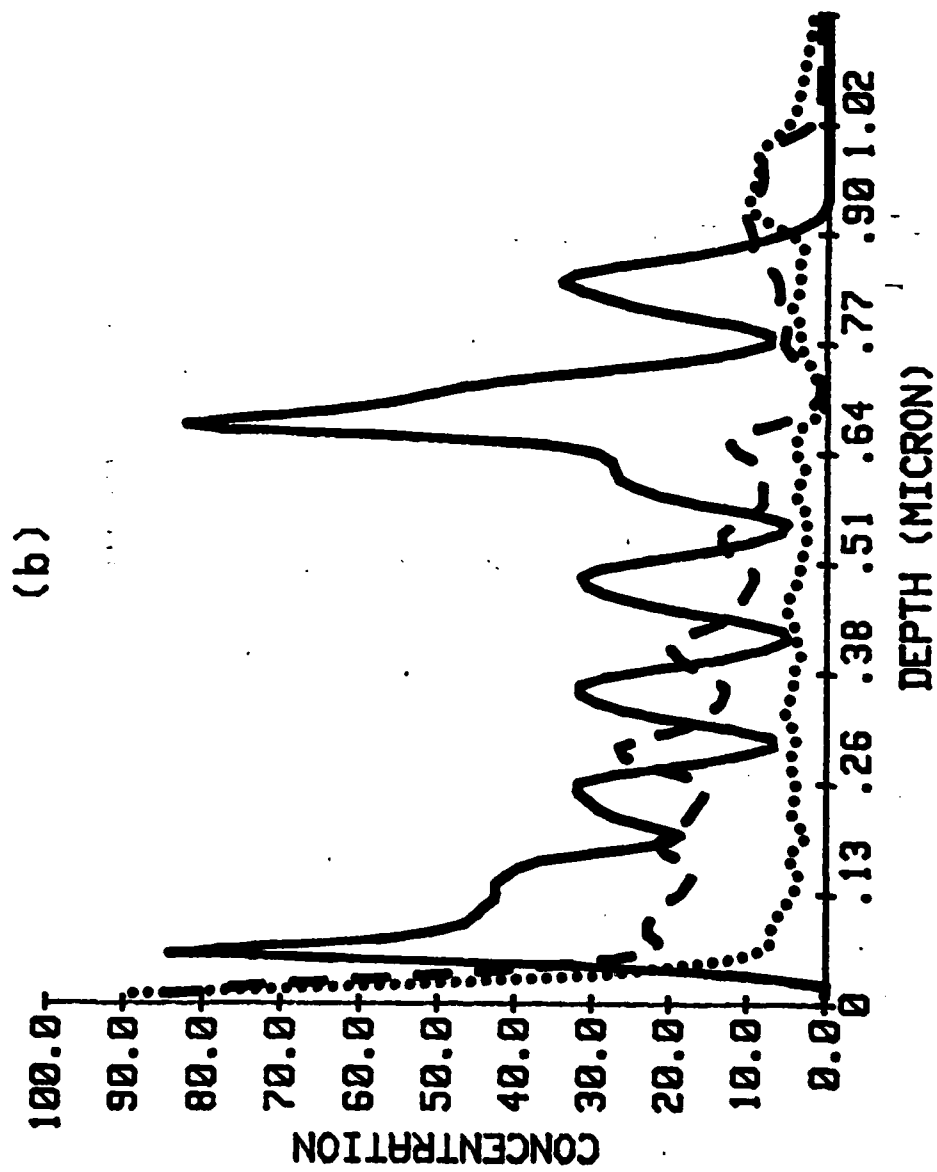












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